## AMENDMENTS TO THE CLAIMS

(Original) A method for determining a bipolar affective disorder or a
predisposition to a bipolar affective disorder, said method comprising detecting a marker that is
linked to map position 4q35.2 of the human genome in a sample derived from a subject, wherein
the detection is indicative of a bipolar affective disorder or a predisposition to a bipolar affective
disorder in the subject.

## (Cancelled)

- (Original) The method according to claim 1 wherein the marker linked to map position 4a35.2 is located within or comprises the FAT gene.
- 4. (Currently amended) A method for determining a bipolar affective disorder or a predisposition to a bipolar affective disorder, said method comprising detecting a marker within a FAT gene or an expression product thereof that is associated with a bipolar affective disorder in a sample derived from a subject, wherein a presence of the marker is indicative of a bipolar affective disorder or a predisposition to a bipolar affective disorder in the subject and wherein the FAT gene comprises a nucleotide sequence selected from the group consisting of:
  - a nucleotide sequence at least 80% identical to the nucleotide sequence set forth in SEQ ID NO: 1;
  - (ii) a nucleotide sequence that encodes a mRNA at least 80% identical to the nucleotide sequence set forth in SEQ ID NO: 2 or 4; and
  - (iii) a nucleotide sequence that encodes a polypeptide comprising an amino acid sequence at least 80% identical to the amino acid sequence set forth in SEQ ID NO: 3 or 5.
  - 5. (Cancelled)
- 6. (Currently amended) The method according to claim 4 wherein the marker is located within the 3' region of the FAT gene that comprises a nucleotide sequence corresponding to the region spanning from nucleotide position 139,260 to nucleotide position 170,001 of SEQ ID NO: 1.
  - (Cancelled)
  - 8. (Cancelled)
  - (Cancelled)
  - (Cancelled)

- 11. (Cancelled)
- 12. (Cancelled)
- (Currently amended) The method according to claim 4+2 wherein the marker comprises polymorphism is a single nucleotide polymorphism (SNP).
  - 14. (Cancelled)
  - (Cancelled)
  - (Cancelled)
  - 17. (Cancelled)
- 18. (Original) The method according to claim 4 wherein the marker comprises a nucleic acid comprising a nucleotide sequence at least about 80% identical to at least about 20 contiguous nucleotides in a sequence selected from the group consisting of:
  - (i) a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2 and SEQ ID NO: 4;
  - (ii) a sequence capable of encoding a polypeptide comprising an amino acid sequence at least 80% homologous to the sequence set forth in SEQ ID NO: 3 and SEQ ID NO: 5; and
    - (iii) a sequence complementary to a sequence set forth in (i) or (ii).
- 19. (Currently amended) The method according to claim 4 wherein the marker is detected by <a href="https://hybridising.nlm.nucleic.acid">hybridising.nucleic.acid</a> in a biological sample derived from a subject under moderate to high stringency <a href="https://hybridisation.nucleic.nucleic.acid">hybridisation.nucleic.nucle
- 20. (Currently amended) The method according to claim 4 wherein the marker is detected by <u>hybridizing</u> <u>hybridising</u> a nucleic acid probe or primer comprising the sequence of the marker to a nucleic acid that is linked to the marker in nucleic acid in a biological sample derived from a subject under moderate to high stringency <u>hybridization</u> <u>hybridisation</u> conditions and detecting the <u>hybridization</u> <u>hybridisation</u> by a detection means, wherein <u>hybridization</u> <u>hybridisation</u> of the probe to the sample nucleic acid indicates that the subject being tested is predisposed to or suffers from a bipolar affective disorder.

21. (Currently amended) The method according to claim 19 o<del>r 20</del>-wherein the detection means is a nucleic acid <u>hybridization</u> <u>hybridisation</u> reaction or a nucleic acid amplification reaction.

- 22. (Cancelled)
- 23. (Cancelled)
- 24. (Original) The method according to claim 4 wherein the marker is detected by contacting a biological sample derived from the subject with an antibody capable of specifically binding to said marker for a time and under conditions sufficient for an antibody-ligand complex to form and then detecting the complex wherein detection of the complex indicates that the subject being tested is predisposed to or suffers from a bipolar affective disorder.
- (Original) The method according to claim 4 wherein the biological sample comprises a nucleated cell.
  - 26. (Cancelled)
  - 27. (Cancelled)
  - 28. (Cancelled)
  - 29. (Cancelled)
  - (Cancelled)
  - (Cancelled)
  - (Cancelled)
  - (Cancelled)
- 34. (Original) A probe or primer comprising at least about 20 nucleotides that is capable of selectively hybridizing to the sequence set forth in SEQ ID NO: 1 and detecting a marker in a FAT gene that is associated with a bipolar affective disorder or a predisposition to a bipolar affective disorder.
  - 35. (Cancelled)
- 36. (Original) A method for determining a subject that carries a gene or allele of a gene or a polymorphism that is associated with a bipolar affective disorder comprising detecting a marker within a FAT gene that is associated with a bipolar affective disorder in a sample derived from the subject, wherein detection of said marker indicates that the subject is a carrier of a gene or allele of a gene or a polymorphism is associated with a bipolar affective disorder.

37. (Previously presented) A method of treatment or prophylaxis of a bipolar affective disorder comprising:

- performing the method of claim 1 for determining a bipolar affective disorder or a predisposition to a bipolar affective disorder; and
- (ii) administering or recommending a therapeutic for the treatment of bipolar affective disorder.
- 38. (Original) A method for identifying a marker that is associated with a bipolar affective disorder, said method comprising:
  - identifying a polymorphism or allele within a FAT gene or an expression product thereof;
  - (ii) analyzing a panel of subjects to determine those that suffer from a bipolar affective disorder, wherein not all members of the panel comprise the polymorphism or allele: and
  - (iii) determining the variation in the development of a bipolar affective disorder wherein said variation indicates that the polymorphism or allele is associated with a subject's predisposition to a bipolar affective disorder.
- (Original) A method for determining a candidate compound for the treatment of a bipolar affective disorder comprising:
  - (i) administering a candidate compound to an animal or cell comprising or expressing a marker within a FAT gene that is associated with a bipolar affective disorder and determining the level of FAT expression in said cell or animal;
  - (ii) administering a candidate compound to an animal or cell that does not comprise or express a marker within a FAT gene that is associated with a bipolar affective disorder and determining the level of FAT expression in said cell or animal; and
    - (iii) comparing the level of FAT expression at (i) and (ii),

wherein a decreased level of FAT expression at (i) relative to (ii) indicates that the compound is a candidate compound for the treatment of a bipolar affective disorder.

40. (Original) A method for determining a candidate compound for the treatment of a bipolar affective disorder comprising:

> (i) administering a candidate compound to an animal or cell capable of expressing a FAT gene and determining the level of FAT expression in said cell or animal:

- (ii) determining the level of FAT expression in an animal or cell capable of expressing a FAT gene in the absence of the candidate compound; and
  - (iii) comparing the level of FAT expression at (i) and (ii),

wherein a decreased level of FAT expression at (i) relative to (ii) indicates that the compound is a candidate compound for the treatment of a bipolar affective disorder.

## 41. (Cancelled)

- 42. (Currently amended) A process for identifying or determining a compound or modulator for the treatment of a bipolar affective disorder said method comprising:
  - (i) performing the method according to either claim 39 er-40 to thereby identify or determine a compound for the treatment of a bipolar affective disorder;
    - (ii) optionally, determining the structure of the compound;
    - (iii) optionally, providing the name or structure of the compound; and
    - (iv) providing the compound.
- 43. (Currently amended) A process of manufacturing a compound for the treatment of a bipolar affective disorder comprising:
  - determining a candidate compound for the treatment of a bipolar affective disorder by performing the method according to either claim 39 or 40;
  - (ii) using the compound in the manufacture of a therapeutic or prophylactic for the treatment of bipolar affective disorder.